

Amended claims

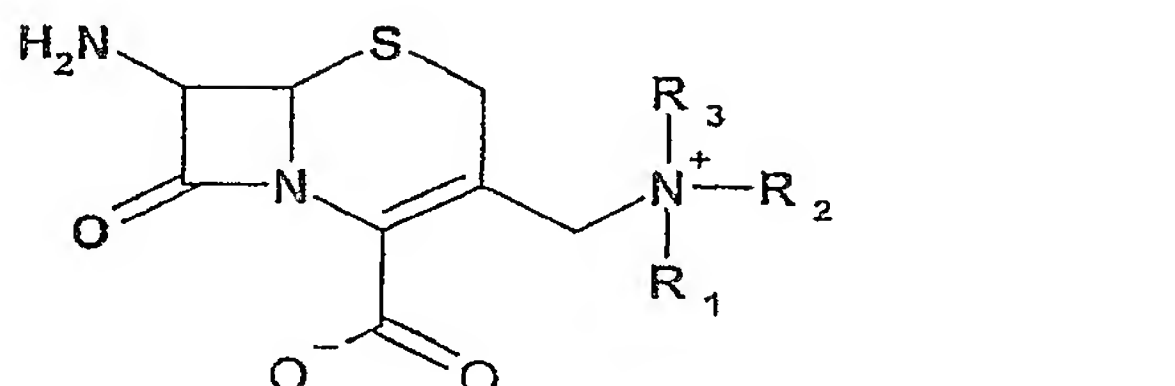
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What we claim is

1. Process for the production of a compound of formula

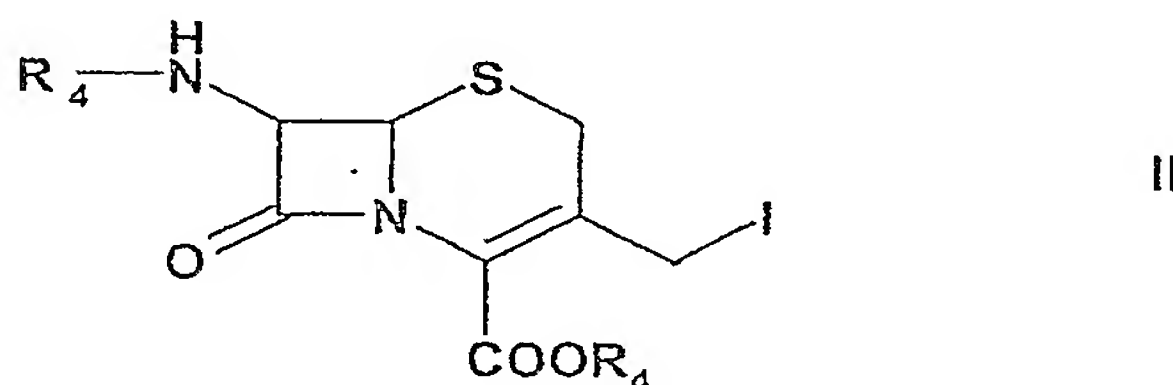


wherein  $R_1$ ,  $R_2$  and  $R_3$ , independently of one another, are alkyl, alkenyl, aryl, hydroxy-( $C_{1-6}$ )alkyl, carbamoyl-( $C_{1-6}$ )alkyl, amino-( $C_{1-6}$ )alkyl, acylamino-( $C_{1-6}$ )alkyl or carboxy-( $C_{1-6}$ )alkyl, or wherein

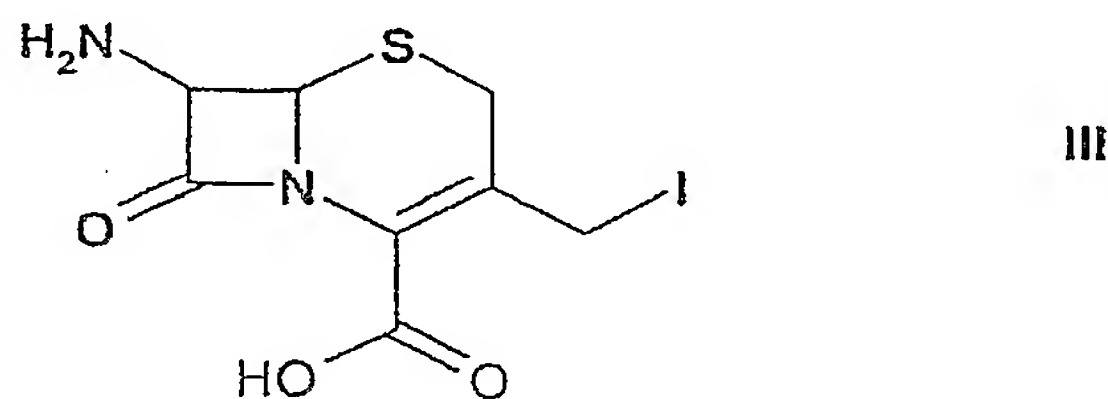
$R_2$  and  $R_3$  together with the adjacent nitrogen atom, form an alicyclic 5- to 8-membered heterocyclic ring, which, in addition to the nitrogen atom, may also contain a further 1 or 2 hetero atoms selected from the group consisting of oxygen and sulphur, and  $R_1$  signifies alkyl, alkenyl or aryl, as well as for the production of acid addition salts and/or hydrates of a compound of formula I,

comprising the reaction steps

- a) desilylation of a compound of formula



wherein  $R_4$  is a silyl-protecting group, by adding a protic solvent, in order to obtain a compound of formula

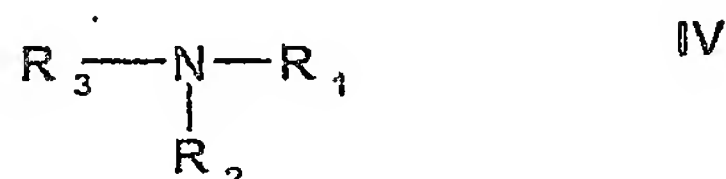


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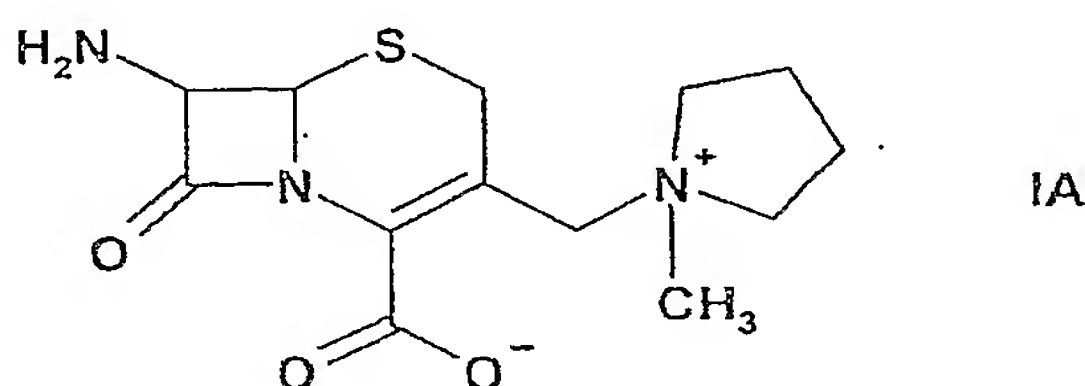
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b) reaction of the compound of formula III obtained in step a) with an organic base of formula



wherein R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub> have the significances indicated above, in order to obtain a compound of formula I, and

- ~~2.~~ Process according to ~~claim 1~~, whereby steps a) and b) are carried out simultaneously in one reaction container.
- ~~2.~~ <sup>3</sup> Process according to ~~one of claims 1, or 2~~ whereby R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub>, independently of one another, are alkyl, alkenyl, aryl or hydroxy(C<sub>1-6</sub>)alkyl.
- ~~3.~~ <sup>4</sup> Process according to ~~one of claims 1 or 2~~, whereby R<sub>2</sub> and R<sub>3</sub> together represent a C<sub>4</sub>-alkylene group, and with the adjacent nitrogen atom, form a saturated 5-membered heterocycle, and R<sub>1</sub> represents a methyl group, so that a compound of formula



is obtained.

- ~~4.~~ <sup>3</sup> Process according to one of claims 1 to ~~4~~, wherein the protic solvent is a (C<sub>1-4</sub>)-alcohol or a mixture of several (C<sub>1-4</sub>)-alcohols.
- ~~5.~~ <sup>4</sup> Process according to claim ~~5~~, wherein the alcohol is methanol, ethanol, isopropanol, n-propanol, 2-methyl-propan-2-ol, glycol, glycerol, a propanediol or a butanediol.
- ~~6.~~ <sup>3</sup> Process according to claim ~~6~~, whereby the alcohol is isopropanol or 1,2-butanediol.

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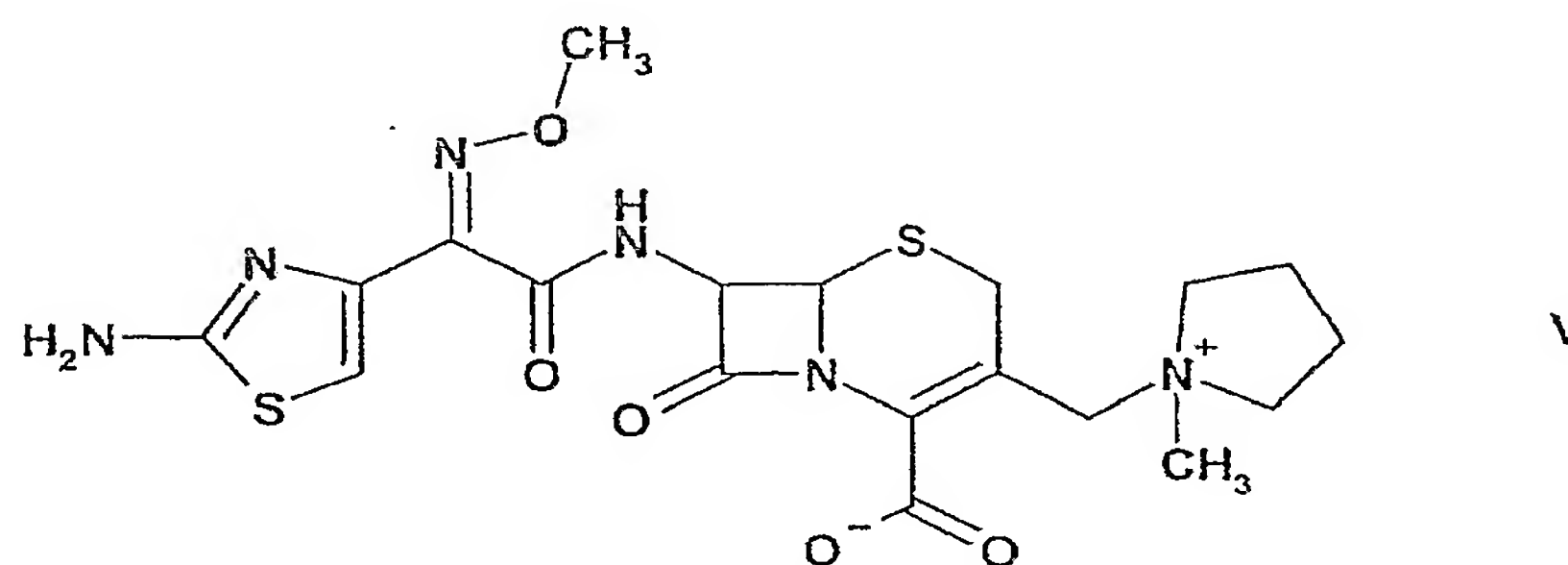
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~~7~~ ~~8~~. Process according to one of the preceding claims, whereby a compound of formula I obtained from step b) is obtained in the form of an acid addition salt and/or hydrate or is converted into the same.

~~8~~ ~~9~~. Process according to claim ~~8~~ <sup>7</sup>, whereby the acid addition salt is a hydriodide or a hydrochloride.

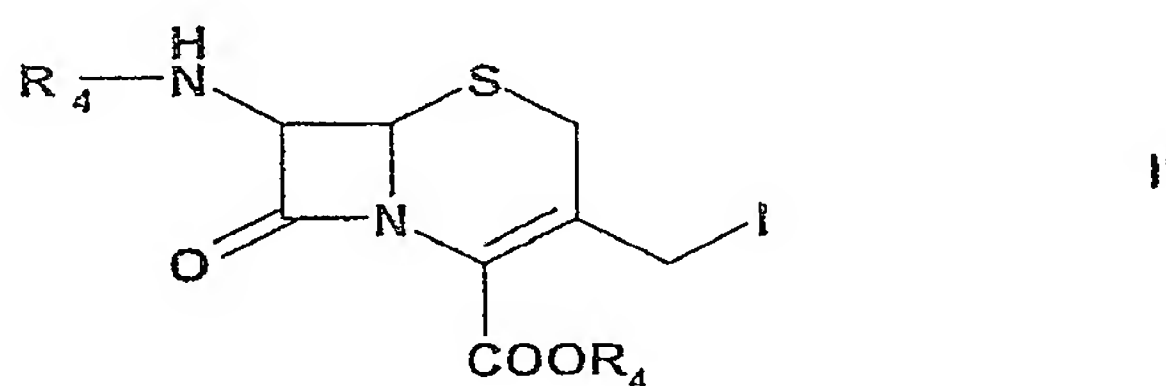
~~9~~ ~~10~~. Process according to one of claims ~~8 to 9~~ <sup>7 to 8</sup>, whereby the hydrate is a monohydrate.

~~10~~ ~~11~~. Process for the production of cefepime of formula

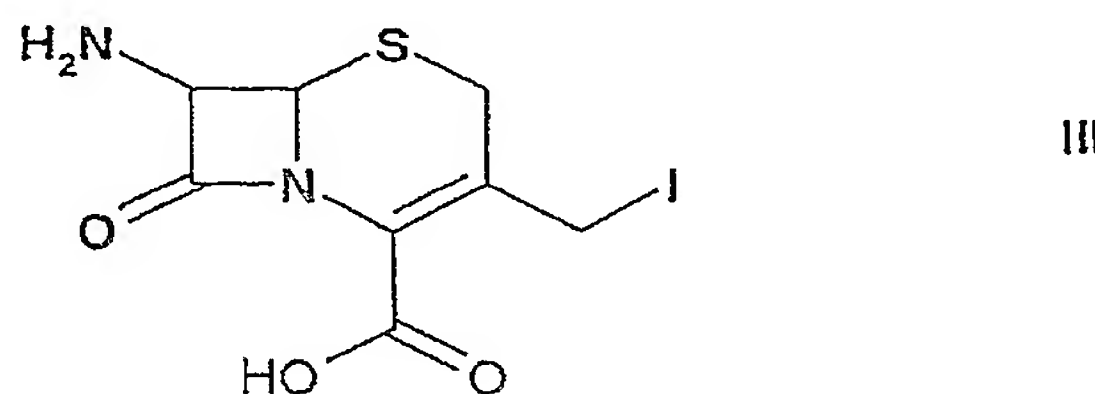


or one of its acid addition salts and/or its hydrates comprising the reaction steps

a) desilylation of a compound of formula



wherein R<sub>4</sub> is a silyl-protecting group, by adding a protic solvent, in order to obtain a compound of formula

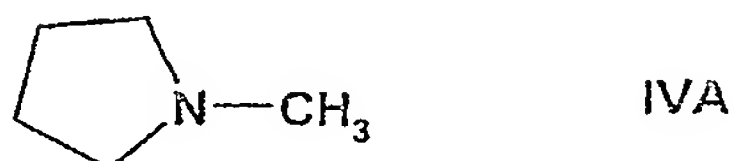


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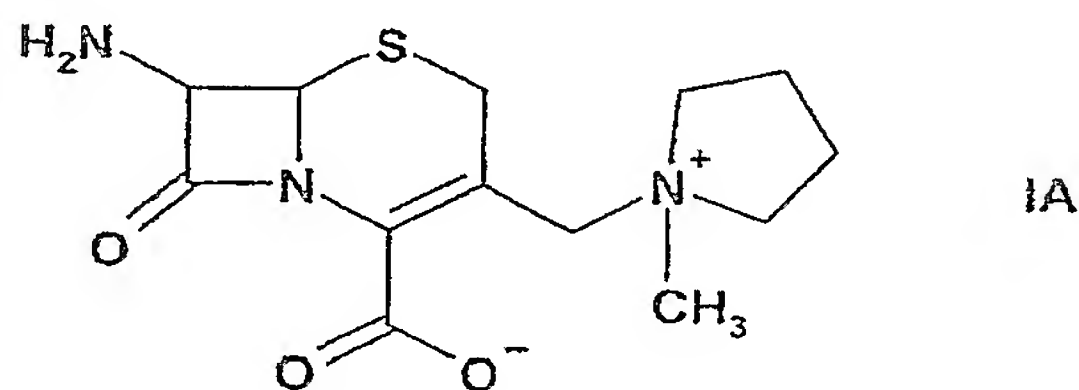
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b) reaction of the compound of formula III obtained in step a) with a strong organic base of formula



in order to obtain a compound of formula



- c) optional conversion of a compound of formula IA, as obtained from step b), into the form of an acid addition salt and/or a hydrate, and
- d) acylation of the 7-amino group of a compound of formula IA obtained from step b) or of its acid addition salt and/or hydrate obtained from step c), in order to obtain cefepime of formula V,

whereby steps a) and b) are carried out simultaneously in one reaction container.